Belgian Genetic Tests database

ANALYTE:
NOTCH1

NAME:	notch receptor 1
SYMBOL:	NOTCH1
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Adams-Oliver syndrome (gene panel)
- Aneurysm, Thoracic Aortic, familial (gene panel)
- Bicuspid aortic valve
- Congenital malformation (gene panel - 1721 genes)
- Congenital structural heart defects (gene panel)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Intellectual disability (gene panel)
- Maffucci syndrome (gene panel)
- Overgrowth & vascular anomalies (gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skin disorders (gene panel)
- Sturge-Weber syndrome (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Adams-Oliver syndrome
- Familial bicuspid aortic valve

Related Gene Panels

- Adams-Oliver (6 genes) - UZA

- Bicuspid aortic valve - UGent
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital structural heart defects - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Familial Thoracic Aortic Aneurysm (genepanel) - UZA
- Heterotaxie PCD - UGent
- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Maffucci syndrome (65 genes) - KUL
- Overgrowth & vascular anomalies (65 genes) - KUL
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent
- Sturge-Weber syndrome (65 genes) - KUL

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